Under the Paperwork Reduction Act of 1995, no persons are required to respond to

publisher, city and/or country where published.

INFORMATION DISCLOSURE
STATEMENT BY APPLICANT
STATEMENT BY APPLICANT
(Not for submission under 37 CFR 1.99)

Application Number		10596507		
iling Date				
First Named Inventor Danie		elle Louise Lehrer		
Art Unit				
Examiner Name				
Attorney Docket Number		LEHR0101PUSA	Ī	

					U.S.	PATENTS			Remove		
Examiner Initial*	Cite No	Patent Number	Kind Code ¹	Issue E	Date	Name of Patentee or Applicant of cited Document		Releva	Columns,L int Passage s Appear		
	1	6510236	B1	2003-0	2-21	Crane et al					
If you wish	h to a	dd additional U.S. Pate	nt citatio	n inform	ation pl	ease click the	Add button.		Add		
			U.S.P	ATENT	APPLI	CATION PUB	LICATIONS		Remove		
Examiner Initial*	Cite No	Publication Number	Kind Code ¹	Publica Date	tion	Name of Patentee or Applicant of cited Document		Releva	Columns,L int Passage s Appear		
	1	20020188736	A1	2002-1	2-12	Jarvensivu					
If you wish	h to a	dd additional U.S. Publi	shed Ap	plication	citatio	n information p	olease click the Ad	d button	Add		
				FOREIG	SN PAT	TENT DOCUM	IENTS		Remove		
Examiner Initial*	Cite No	Foreign Document Number ³			Kind Code ⁴	Publication Date	Name of Patentee or Applicant of cited Document		Pages,Colu vhere Relev Passages o Figures App	/ant r Relevant	+-
	1	2000035143	wo		A1	2000-06-15	Virtual Business Associates Pty. Lte	1.			
If you wish	h to a	dd additional Foreign P	atent Do	cument	citation	information p	lease click the Add	button	Add		
			NON	-PATE	NT LITE	RATURE DO	CUMENTS		Remove		
Examiner	Cite	Include name of the a									Τs

	Application Number		10596507	
NEODIA TION DIOCI COURT	Filing Date			
INFORMATION DISCLOSURE STATEMENT BY APPLICANT	First Named Inventor Danie		elle Louise Lehrer	
(Not for submission under 37 CFR 1.99)	Art Unit			
	Examiner Name			
	Attorney Docket Number		LEHR0101PUSA	

	1			
If you wis	h to a	id additional non-patent literature document citation information please click the Add button	Add	

you wish to add additional non-patient illerature document cliation information please click the Add Button | Your |

EXAMINER SIGNATURE

Examiner Signature

Date Considered

*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through a citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

See Kint Code of USPTO Patent Documents at invenUSPTO_GDI/ or MPEP 901.04. * Enter of thice that issued the document, by the involved ow (WIPO Standard ST3.). * For Suparese patent counters, the noticeation of the year of the region of the Emperor many precede the sent annual replication of the parent of the patent document. * Varied of counters by the appropriate symbols as endicated on the document under WIPO Standard ST1.6 if possole, * Applicant is to place a check mark here if Emploit languages translation is attached.

Application Number | 10596507 | Filing Date |

CERTIFICATION STATEMENT

That each item of information contained in the information disclosure statement was first cited in any communication from a foreign patient office in a counterpart foreign application not more than three months prior to the filing of the information disclosure statement. See 37 CFR 1.97(e)(1).

OR

That no item of information contained in the information disclosure statement was cited in a communication from a foreign patent office in a counterpart foreign application, and, to the knowledge of the person signing the certification after making reasonable inquiry, no item of information contained in the information disclosure statement was known to any individual designated in 37 CFR 1.59(c) more than three months prior to the filing of the information disclosure statement. See 37 CFR 1.97(c) and the statement. See 37 CFR 1.97(c) and the statement. See 37 CFR 1.97(c) and the statement. See

- See attached certification statement.
- Fee set forth in 37 CFR 1.17 (p) has been submitted herewith.

Please see 37 CFR 1 97 and 1 98 to make the appropriate selection(s):

.7 None

SIGNATURE

A signature of the applicant or representative is required in accordance with CFR 1.33, 10.18. Please see CFR 1.4(d) for the form of the signature.

of the signature.							
	Signature	/John E, Nemazı/	Date (YYYY-MM-DD)	2006-08-18			
	Name/Print	John Nemazi	Registration Number	30876			

This collection of information is required by 37 CFR 1.97 and 1.98. The information is required to obtain or retain a benefit by the public which is for life and by the USPTO to process) an application. Confidentiality is governed by \$5 U.S. C.12 and 37 CFR 1.14. This collection is estimated to take 1 hour to complete, including gathering, preparing and submitting the completed application from the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patient and Tradenant Afford, U.S. Department of Commence, P. 0. Bast 1436, Alexandria, V.32.11.4450 D.O. NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, V.32.211.4450

Privacy Act Statement

The Privacy Act of 1974 (P.L. 93-579) requires that you be given certain information in connection with your submission of the attached form related to a patient application or patient. Accordingly, pursuant to the requirements of the Act, please be advised that. (1) the general authority for the collection of this information is 35 U.S.C. 2(b)(2); (2) famishing of the information solicided is voluntary, and (3) the principal purpose for which the information is used by the U.S. Patient and Trademan Kolfice is to process another examine your submission relation to a patient application or patient. If you do not furnish the requested process another examine your submission relation to the patient application or patient. If you do not furnish the requested the process another examines your submission, which may visually intermediate or for extension or about those when the basic high process another examines your submission, which may visually intermediate or for extension or a submission of the basic high process another examines your submission, which may

The information provided by you in this form will be subject to the following routine uses:

- The information on this form will be treated confidentially to the extent allowed under the Freedom of Information Act (5 U.S.C. 552) and the Privacy Act (5 U.S.C. 552a). Records from this system of records may be disclosed to the Department of Justice to determine whether the Freedom of Information Act requires disclosure of these record s.
 - A record from this system of records may be disclosed, as a routine use, in the course of presenting evidence to a court, magistrate, or administrative tribunal, including disclosures to opposing counsel in the course of settlement neodiations.
 - A record in this system of records may be disclosed, as a routine use, to a Member of Congress submitting a request involving an individual, to whom the record partains, when the individual has requested assistance from the Member with respect to the subject matter of the record.
 - A record in this system of records may be disclosed, as a routine use, to a contractor of the Agency having need for the information in order to perform a contract. Recipients of information shall be required to comply with the requirements of the Privacy Act of 1974. as amended, pusuant to 5 U.S.C. 552a(m).
 - A record related to an International Application filed under the Patent Cooperation Treaty in this system of records
 may be disclosed, as a routine use, to the International Bureau of the World Intellectual Property Organization, pursuant
 to the Patent Cooperation Treaty.
 - A record in this system of records may be disclosed, as a routine use, to another federal agency for purposes of National Security review (35 U.S.C. 181) and for review pursuant to the Atomic Energy Act (42 U.S.C. 218(c)).
- 7. A record from this system of records may be disclosed, as a routine use, to the Administrator, General Services, or hisher designe, cuting an inspection of records concluded by GSAs a part of that apency's responsibility to recommend improvements in records management practices and programs, under suthority of 4d U.S.C. 2004 and 2006. Such disclosure shall be made in accordance with the GSA regulations governing inspection of records for this purpose, and any other relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make determinations about individuals.
- A record from this system of records may be disclosed, as a routine use, to the public after either publication of the application pursuant to 35 U.S. C. 122(b) or issuance of a patent pursuant to 35 U.S. C. 151. Further, a record may be disclosed, subject to the limitations of 37 CFR 1.14, as a routine use, to the public if the record via set float in an application which became abandomed or in which the proceedings were terminated and which application is referenced by either a published application, an application open to public inspections or an issuand patent.
 - A record from this system of records may be disclosed, as a routine use, to a Federal, State, or local law enforcement agency, if the USPTO becomes aware of a violation or potential violation of law or regulation.